

KOS Diagnostic Lab

(A Unit of KOS Healthcare)



Dr. Vinay Chopra MD (Pathology & Microbiology) Chairman & Consultant Pathologist Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist

NAME: Miss. KHUSHANI 113152 Accession No.: Age/Gender: 8 Y/Female Specimen ID: BC2401168 012412050011 Whole Blood HEPARIN Lab NO: Specimen: Referred BY: Self Collected: 05/Dec/2024 01:35PM Registered: 05/Dec/2024 01:32PM Remark: Reported: 24/Dec/2024 05:29PM

CYTOGENETICS REPORT

Test Name:Karyotype - Blood

RESULT:

 Method:
 G-banding

 Metaphases counted:
 20

 Metaphases analyzed:
 20

 Metaphases karyotyped:
 14

 Banding Resolution:
 400

 Karyotype (ISCN 2016):
 46,XX

INTERPRETATION:

Normal female chromosome complement. There is no evidence of aneuploidy or structural rearrangement at the resolution of banding analysis.

RECOMMENDATIONS:

Chromosome microarray analysis is recommended for this patient because this test will be able to detect submicroscopic deletions and duplications in the genome, which cannot be detected by chromosome analysis. CMA is now considered the first-tier cytogenetic diagnostic test (Miller et al., 2010; Manning, Hudgins and the ACMG Professional Practice and Guidelines Committee, 2010). This testing is now available in our Laboratory, contact us for more information. In addition, a complete genetic evaluation should be considered to rule out other genetic etiologies associated with the clinical finding(s) in this patient. Genetic counseling is recommended.



Mr. Brijesh Authorised Signatory PhD(P)

DR. S. KUMAR MBBS, MD Consultant Pathologist



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NOTE:



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CYTOGENETICS REPORT

KARYOTYPE:

2 3 4 5 5 5 6 7 8 0 10 11 12 12 13 14 15 16 17 18

Disclaimer: Although the methodology used in this analysis and interpretation is highly accurate, it does not detect small rearrangements and very low-level mosaicism, which are detectable only by molecular methods. Failure to detect an alteration at any locus does not exclude the diagnosis of any of the disorders. LABASSURE can assist the physician in determining the appropriate test in the context of clinical indications.

*** End Of Report ***







